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I hereby request access under 37 CFR 1.14(a)(3)(iv) to the application file record of the above-identified ABANDONED application, which is: (CHECK ONE)

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United States Patent [19]

Queen et al.

[11] Patent Number: 5,585,089

[45] Date of Patent: Dec. 17, 1996

[54] HUMANIZED IMMUNOGLOBULINS

[75] Inventors: Cary L. Queen, Los Altos; Harold E. Selick, Belmont, both of Calif.

[73] Assignee: Protein Design Labs, Inc., Mountain View, Calif.

[21] Appl. No.: 477,728

[22] Filed: Jun. 7, 1995

Related U.S. Application Data

[63] Continuation of Ser. No. 634,278, Dec. 19, 1990, Pat. No. 5,530,101, which is a continuation-in-part of Ser. No. 590,274, Sep. 28, 1990, abandoned, and Ser. No. 310,252, Feb. 13, 1989, abandoned, which is a continuation-in-part of Ser. No. 290,975, Dec. 28, 1988, abandoned.

[51] Int. Cl.⁶ C07K 16/18; A61K 39/395

[52] U.S. Cl. 424/133.1; 530/387.3; 530/388.22; 424/143.1

[58] Field of Search 530/387.3, 388.22; 424/133.1, 143.1

[56] References Cited

U.S. PATENT DOCUMENTS

4,578,335	3/1986	Urdal et al.	530/351
4,816,397	3/1989	Boss et al.	435/68
4,816,565	3/1989	Honjo et al.	435/69.1
4,816,567	3/1989	Cabilly et al.	530/387
4,845,198	7/1989	Urdal et al.	530/387.3
4,867,973	9/1989	Goers et al.	
5,198,359	3/1993	Taniguchi et al.	435/252.3
5,225,539	7/1993	Winter	530/387.3

FOREIGN PATENT DOCUMENTS

0171496	2/1986	European Pat. Off.	C12N 15/00
0173494	3/1986	European Pat. Off.	C12N 15/00
0184187	6/1986	European Pat. Off.	C12N 15/00
0256654	7/1987	European Pat. Off.	
0239400	9/1987	European Pat. Off.	
0266663	6/1988	European Pat. Off.	C12N 15/00
2188941	10/1987	United Kingdom	C12N 5/00
86/05513	9/1986	WIPO	C12N 15/00
87/02671	5/1987	WIPO	C07H 15/12
89/01783	3/1989	WIPO	A61K 39/395

OTHER PUBLICATIONS

Riechmann et al. *Nature* vol. 332 24, Mar. 1988 p. 323.
Footc, *Nova Acta Leopoldina* 1989. vol. 61 (269) 103.
Amit et al. *Science* vol. 233 1986 p. 747.
Groves et al. vol. 6, 1987, p. 71.
Better et al., "Escherichia coli Secretion of an Active Chimeric Antibody Fragment", *Science* 240:1041-1043 (1988).
Bird et al., "Single-Chain Antigen-Binding Proteins", *Science* 242:423-426 (1988).
Boulianne et al., "Production of functional chimeric mouse/human antibody," *Nature* 312:643-646 (1984).
Carter et al., "Humanization of an anti-p185^{HER2} antibody for human cancer therapy," *Proc. Natl. Acad. Sci.* 89:4285-4289 (1992).
Chothia, C. and A. M. Lesk, "Canonical Structures for the Hypervariable Regions of Immunoglobulins", *J. Mol. Biol.* 196:901-917 (1987).

Co et al., "Humanized antibodies for antiviral therapy," *Proc. Natl. Acad. Sci. USA* 88:2869-2873 (1991).

Co et al., "Chimeric and Humanized Antibodies with Specificity for the CD33 Antigen," *J. of Immunol.* 148(4):1149-1154 (1992).

Daugherty et al., "Polymerase chain reaction facilitates the cloning, CDR-grafting, and rapid expression of a murine monoclonal antibody directed against the CD18 component of leukocyte integrins," *Nuc. Acids Res.* 19:2471-2476 (1991).

Ellison et al., "The nucleotide sequence of a human immunoglobulin C(gamma)₂ gene", *Nucleic Acids Res.* 10:4071-(1982).

Farrar, J., "The biochemistry, biology, and role of interleukin-2 in the induction of cytotoxic T cell and antibody-forming B cell receptors," *Immunol. Rev.* 63:129-166 (1982).

Foote et al., "Antibody framework residues affecting the conformation of hypervariable loops," *J. Mol. Biol.* 224:487-499 (1992).

Gorman et al., "Reshaping a therapeutic CD4 antibody," *Proc. Natl. Acad. Sci.* 88:4181-4185 (1991).

Greene et al., "Growth of Human T Lymphocytes: An Analysis of Interleukin 2 and Its Cellular receptor", in *Progress in Hematology XIV*, E. Brown, ed., Grunc and Statton, New York (1986) pp. 283-301.

Hale et al., "Remission Induction in Non-Hodgkin Lymphoma with Reshaped Human Monoclonal Antibody CAMPATH-1H", *Lancet* Dec. 17, 1988, pp. 1394-1399.

Hietter et al., "Cloned Human and Mouse Kappa Immunoglobulin Constant and J Region Genes Conserve Homology in Functional Segments", *Cell* 22:197-207 (1980).

(List continued on next page.)

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[57] ABSTRACT

Novel methods for producing, and compositions of humanized immunoglobulins having one or more complementarity determining regions (CDR's) and possible additional amino acids from a donor immunoglobulin and a framework region from an accepting human immunoglobulin are provided. Each humanized immunoglobulin chain will usually comprise, in addition to the CDR's, amino acids from the donor immunoglobulin framework that are, e.g., capable of interacting with the CDR's to effect binding affinity, such as one or more amino acids which are immediately adjacent to a CDR in the donor immunoglobulin or those within about 3 Å as predicted by molecular modeling. The heavy and light chains may each be designed by using any one or all of various position criteria. When combined into an intact antibody, the humanized immunoglobulins of the present invention will be substantially non-immunogenic in humans and retain substantially the same affinity as the donor immunoglobulin to the antigen, such as a protein or other compound containing an epitope.